What is claimed is:

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- 1. A method for reducing the concentration of an analyte in a blood cell suspension, the method comprising:
- (i) providing a starting blood cell suspension in a volume greater than 50 mL, the blood cell suspension comprising blood cells and extracellular fluid; and
 - (ii) washing the starting blood cell suspension with a wash solution under conditions sufficient to lower the concentration of the analyte at least 10³-fold relative to the analyte concentration in the starting blood cell suspension, wherein the blood cells of the blood cell suspension retain viability after a storage period of greater than 21 days at 4 °C in a storage solution.
 - 2. The method of claim 1, wherein the washing comprises
 - (i) centrifuging the starting blood cell composition to form a pelleted cell fraction and a supernatant;
 - (ii) removing the supernatant from the pelleted cell fraction;
 - (iii) adding washing solution to the pelleted cell fraction; and.
 - (iv) resuspending the pelleted cell fraction in the washing solution to form a resuspended cell suspension;
 - (v) optionally repeating steps (i) (iv); and
 - (vi) resuspending the pelleted cell fraction in a storage solution.
 - 3. The method of claim 2, wherein the analyte is a small molecule.
- 4. The method of claim 3, wherein the small molecule is an ethyleneimine oligomer,
- 25 phenothiazine derivative, acridine derivative, psoralen derivative or riboflavin.
 - 5. The method of claim 3, wherein the small molecule is a therapeutic agent.
 - 6. The method of claim 2, wherein the analyte is a protein.

- 7. The method of claim 6, wherein the protein is a prion protein.
- 8. The method of claim 7, wherein the prion protein is a pathogenic protein.
- 5 9. The method of claim 2, wherein the analyte is a cell.
 - 10. The method of claim 9, wherein the cell is a leukocyte.
- 11. The method of claim 10, wherein the method further comprises treating the starting blood cell10 suspension with an anti-pathogenic agent.
 - 12. The method of claim 11, wherein the anti-pathogenic agent is an ethyleneimine oligomer, phenothiazine derivative, acridine derivative, psoralen derivative or riboflavin.

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